



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁵ : C07C 50/18, 50/34, A61K 31/03 A61K 31/09, 31/12, 31/135</p>	A1	<p>(11) International Publication Number: WO 90/08759 (43) International Publication Date: 9 August 1990 (09.08.90)</p>
<p>(21) International Application Number: PCT/US90/00487 (22) International Filing Date: 26 January 1990 (26.01.90) (30) Priority data: 302,836 27 January 1989 (27.01.89) US (60) Parent Application or Grant (63) Related by Continuation US 302,836 (CIP) Filed on 27 January 1989 (27.01.89) (71) Applicant (for all designated States except US): SLOAN-KETTERING INSTITUTE FOR CANCER RESEARCH [US/US]; 410 East 62nd Street, New York, NY 10021 (US).</p>		<p>(72) Inventors; and (75) Inventors/Applicants (for US only): WATANABE, Kyoichi, A. [JP/US]; 28 Wilton Road, Rye-Brook, NY 10573 (US). KOYAMA, Masao [JP/JP]; 509-3, Minowa-cho, Kouhoku-ku, Yokohama-shi, Kanagawa 223 (JP). (74) Agent: WHITE, John, P.; Cooper & Dunham, 30 Rockefeller Plaza, New York, NY 10112 (US). (81) Designated States: AT (European patent), BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent), US. Published With international search report.</p>
<p>(54) Title: DERIVATIVES OF CHRYSOPHANOL</p> <div style="text-align: center; margin: 20px 0;"> <p>(I)</p> </div> <p>(57) Abstract</p> <p>The present invention concerns compounds of formula (I) wherein R¹ is hydrogen, a hydroxy group or a methoxy group; R² is hydrogen or a methyl group; R³ is hydrogen or a methyl group; Y is a secondary amino group (NHalkyl) or a tertiary amino group (N(alkyl)₂); and Z is hydrogen or a halogen. The invention further concerns pharmaceutical compositions which comprise the above-identified compound or the acid salts thereof, and the use of the compound or compositions for treating a malignancy in a subject.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	ES	Spain	MG	Madagascar
AU	Australia	FI	Finland	ML	Mali
BB	Barbados	FR	France	MR	Mauritania
BE	Belgium	GA	Gabon	MW	Malawi
BF	Burkina Faso	GB	United Kingdom	NL	Netherlands
BG	Bulgaria	HU	Hungary	NO	Norway
BJ	Benin	IT	Italy	RO	Romania
BR	Brazil	JP	Japan	SD	Sudan
CA	Canada	KP	Democratic People's Republic of Korea	SE	Sweden
CF	Central African Republic	KR	Republic of Korea	SN	Senegal
CG	Congo	LI	Liechtenstein	SU	Soviet Union
CH	Switzerland	LK	Sri Lanka	TD	Chad
CM	Cameroon	LJ	Luxembourg	TG	Togo
DE	Germany, Federal Republic of	MC	Monaco	US	United States of America
DK	Denmark				

DERIVATIVES OF CHRYSOPHANOL

The invention described herein was made in the course of work under Grant Nos. CA-08748 and CA-18856 from the National Cancer Institute, National Institutes of Health, U.S. Department of Health and Human Services. The U.S. Government has certain rights in this invention.

Background of the Invention

This application is a continuation-in-part of U.S. Serial No. 302,836, filed January 27, 1989, the contents of which are hereby incorporated by reference into the present disclosure.

Some of the information set forth herein has been published (see Masao Koyama, T. Ross Kelly and Kyoichi A. Watanabe, Novel Type of Potential Anticancer Agents Derived from Chrysophanol and Emodin. Some Structure-Activity Relationship Studies, [Part 1] Journal of Medicinal Chemistry, 31:283-284 (1988) which was distributed by the publisher on January 29, 1988; and Masao Koyama, Kiyobumi Takahashi, Ting-Chao Chou, Zbigniew Darzynkiewicz, Jan Kapuscinski, T. Ross Kelly, and Kyoichi A. Watanabe, Intercalating Agents with Covalent Bond Forming Capability. A Novel Type of Potential Anticancer Agents. [Part 2] Derivatives of Chrysophanol and Emodin, Journal of Medicinal Chemistry, 32:1594 (1989)).

A number of analogues of certain antitumor intercalating agents, such as ellipticine (Le Pecq, J.-B., et al., Proc. Natl. Acad. Sci. U.S.A. 71:5078 (1974); Guthrie, R. W., et al., J. Med. Chem 18:755 (1975)) 4'-(9-acridinyl-amino)methanesulfon-*m*-aniside (*m*-AMSA, amsacrine), (Denny, W.A., et al., J. Med. Chem. 25:276 (1982)) and anthracycline antibiotics (e.g., doxorubicin) (Mosher, C.W., et al., J.

Med. Chem. 25:18 (1982); Seshadri, R., et al., J. Med. Chem. 26:11 (1983); Myers, C. Cancer Chemother. 8:52 (1986)) have been synthesized in order to gain better therapeutic potential. However, preliminary screening data show that there is no straightforward structure-activity relationship within each group. These results seem to suggest that although intercalation may be a necessary condition, it may not be sufficient and other factors may be involved that per se potentiate the anticancer activity.

Studies on the mechanism of anticancer action of antibiotic CC1065 (Chidester, C.G., et al., J. Am. Chem. Soc., 103: 7629 (1981); Kanatomo, S., et al., Chem. Pharm. Bull., 29:229 (1981), Li, L.H. et al., Cancer Res. 42:999 (1982)) show that it binds to the minor groove of DNA by non-intercalative means and then slowly alkylates the amino group of adenine by opening the cyclopropane ring in the antibiotic molecule. With CC1065, covalent binding of the drug with DNA, therefore, seems to be important for its potent cytotoxic activity. Mere physical interaction between the drug and DNA may not be sufficient.

These considerations point to the development of intercalators with slow alkylating capability. Such intercalators will bind covalently and hopefully should eventually disrupt the DNA function.

The compounds of the present invention have both intercalating and alkylating functionalities, and as such are potential anticancer agents.

The compounds of this invention may also be useful as biochemical probes for biological reactions essential for DNA synthesis.

Recent studies indicate that m-AMSA inhibits the topoisomerization and catenation reactions of DNA

topoisomerase II (Wang, J.C. Annu. Rev. Biochem, 54:665 (1985)), probably by trapping the enzyme-DNA complexes. (Nelson, E.M., et al., Proc. Natl. Acad. Sci. U.S.A. 81:1361 (1984)); Chen, G.L., et al., J. Biol. Chem. 259:13560 (1984)). Other substances, such as etoposide (VP-16), adriamycin, and ellipticine (Kuhn, K.W., et al., Natl. Cancer Inst. Monogr. 4:61 (1987)) also stabilize the cleavable complex between DNA topoisomerase II and DNA.

In the present invention, we show that the incorporation of an alkylating group into some DNA intercalating agents greatly enhances their antileukemic properties.

15

20

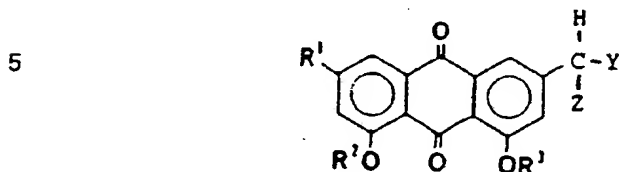
25

30

35

Summary of the Invention

The present invention concerns compounds of the formula:



10

wherein R¹ is hydrogen, a hydroxy group or a methoxy group;
R² is hydrogen or a methyl group;
R³ is hydrogen or a methyl group;
Y is a halogen, secondary amino group or a
15 tertiary amino group and
Z is hydrogen or a halogen.

The invention further concerns pharmaceutical compositions
which comprise the above-identified compound or the acid
20 salts thereof, and the use of the compound or compositions
for treating a malignancy in a subject.

25

30

35

Brief Description of the Figures

Figure 1: Derivatives of chrysophanol and emodin.

5 Figure 2: Visible light absorption spectrum of derivative XIb-6 (11.3 mM in the buffer containing 0.01 M NaCl) alone (solid line) and in the presence of 0.2 mM calf thymus DNA (Sigma type 1) (broken line).

10 Figure 3: Biological effects of chrysophanol derivatives.

Figure 4: Topo II cleavable complex formation. (Compounds SK-31833; SK-31824; SK-31660).

15 Figure 5: Topo II cleavable complex formation. (Compounds SK-31694; SK-31669; SK-31662).

20 Figure 6: Topo II cleavable complex formation. (Compounds SK-31661; SK-31671; SK-31690).

20

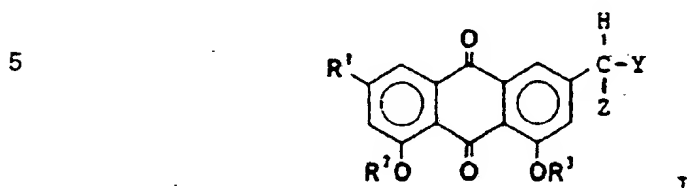
25

30

35

Detailed Description of the Invention

The present invention concerns compounds of the formula (I):



10

wherein R¹ is hydrogen (H), a hydroxy group (OH) or a methoxy group (OMe);

R² is hydrogen or a methyl group (Me);

R³ is hydrogen or a methyl group;

15 Y is a halogen, a secondary amino group or a tertiary amino group and

Z is hydrogen or a halogen.

When Y is a secondary amino group, e.g. [-NHalkyl], or a
20 tertiary amino group, e.g. [-N(alkyl)₂], it is preferred that the alkyl groups be lower alkyl groups, e.g. groups having from one to about five carbon atoms. Particularly effective are methyl or ethyl groups. The lower alkyl groups may also have substituents on the carbon atoms for
25 the hydrogens. The alkyl groups may be substituted with one or more hydroxyl group(s), for example 2-hydroxyethyl, or formed into organic acyl esters, such as acetyl, benzoyl or methanesulfonyl esters, i.e. the substituents are benzoxy, acetyloxy, or methylsulfonyloxy. Further, the alkyl groups
30 may be substituted with halogens(s), such as chlorine and/or bromine to form groups such as a 2-chloroethyl or a 2-bromoethyl.

The invention also provides for pharmaceutical compositions
35 for the treatment of a malignancy in a subject comprising

the compound or a pharmaceutically acceptable acid salt thereof, and a pharmaceutically acceptable carrier, the amount of the composition being an amount effective to suppress the growth of the malignancy, preferably from 1-200 mg/kg of the body weight of the subject.

- 5 The invention further provides for a method of treating a subject have a malignancy which comprises administering to the subject an effective amount of the compound to suppress the growth of the malignancy. A subject may be any warm-blooded animal, preferably human. The malignancy is
10 preferably a tumor or leukemia.

The following Experimental Detail Section and Examples are set forth to aid in an understanding of the invention. These sections are not intended to, and should not be
15 construed to, limit in any way the invention set forth in the claims which follow thereafter.

20

25

30

35

Experimental Detail

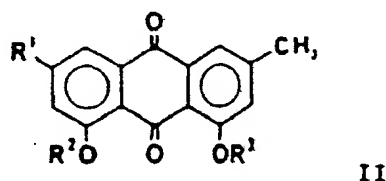
The present invention provides a novel class of compounds, anthraquinones, which possess covalent bonding capability. Such compounds may intercalate into DNA and then bind covalently to DNA, thereby exerting cytotoxic activity.

5

EXPERIMENT 1:

The starting materials for the compounds of the present experiment are of the formula II:

10



15

Typical examples contain the following combinations of R¹, R², and R³.

20

	R ¹	R ² and R ³	
Ila	H	H	chrysophanol
I Ib	OH	H	emodin
I Ic	H	Me	
I Id	OMe	H	
I Ie	OMe	Me	

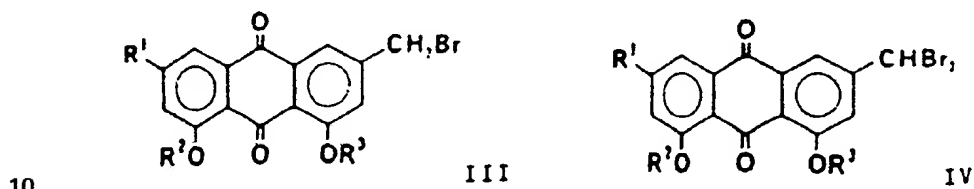
25

30 Compounds I Ic-I Ie are known, and can be prepared readily from the natural product, chrysophanol or emodin (I Ia or I Ib), by the known procedures.

Compounds of formula II are treated with N-bromosuccinimide
 35 (NBS) or 1,3-dibromo-5,5-dimethylhydantoin (BMH) in a

halogenated hydrocarbon, preferably carbon tetrachloride, in the presence of a peroxide, such as m-chloroperbenzoic acid or benzoyl peroxide to give the corresponding monobromides of the formula III as the major products and dibromides of formula IV as the minor products:

5



15 The reaction is carried out at a temperature range of from 25°C to 100°C, preferably at the boiling temperature of the solvent (77°C for carbon tetrachloride) in a period from one hour to three days. The molar ratio of the reactants, formula II to NBS or BMH, can be from 1:1 to 1:3, preferably
20 1:1.2. Upon completion of the reaction, insoluble materials are removed by filtration, the filtrate concentrated, and the residue recrystallized to give formula III compounds. From the mother liquor, IV compounds can be obtained after chromatography on a silica gel column.

25

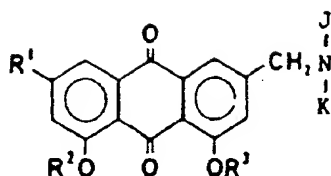
The 1,8-dimethoxy derivatives IIIc, IIIe, IVc and IVe can be converted into the corresponding 1,8-dihydroxy derivatives IIIa, IIId, IVa, and IVd, respectively, by treatment with hydrogen bromide in acetic acid.

30

Treatment of compounds of formula III with a primary or secondary amine with or without solvent affords the corresponding products of formula V:

35

10



5

Some of the typical examples contain the following combinations of R¹, R², R³, J and K.

10

	V	R¹	R² and R³	J	K
15	Va	H	H	Et	Et
	Vb	H	H	CH₂CH₂OH	CH₂CH₂OH
	Vc	H	H	Et	H
	Vd	H	H	CH₂CH₂OH	H
	Ve	OMe	H	Et	Et
20	Vf	OMe	H	CH₂CH₂OH	CH₂CH₂OH
	Vg	OMe	H	Et	H
	Vh	OMe	H	CH₂CH₂OH	H
	Vi	H	Me	Et	Et
	Vj	H	Me	CH₂CH₂OH	CH₂CH₂OH
	Vk	H	Me	Et	H
25	Vl	H	Me	CH₂CH₂OH	H
	Vm	OMe	Me	Et	Et
	Vn	OMe	Me	CH₂CH₂OH	CH₂CH₂OH
	Vo	OMe	Me	Et	H
30	Vp	OMe	Me	CH₂CH₂OH	H

The 1,8-dimethoxy derivatives of formula V (Vi-Vp) can be converted into their corresponding 1,8-

35

dihydroxyanthraquinones of formula V (Va-Vh) by treatment with hydrogen bromide in acetic acid.

The 2-hydroxyethylamino derivatives (Vb, Vd, Vf, Vh, Vj, Vl, Vn and Vp) can be further converted into their corresponding 2-chloroethyl derivatives (Vq-Vx) by treatment with a conventional chlorinating agent, such as thionyl chloride, sulfonyl chloride, phosphorus oxychloride or carbon tetrachloride and triphenyl-phosphine.

V	R ¹	R ² and R ³	J	K
Vq	H	H	CH ₂ CH ₂ Cl	CH ₂ CH ₂ Cl
Vr	H	H	CH ₂ CH ₂ Cl	H
Vs	OMe	H	CH ₂ CH ₂ Cl	CH ₂ CH ₂ Cl
Vt	OMe	H	CH ₂ CH ₂ Cl	H
Vu	H	Me	CH ₂ CH ₂ Cl	CH ₂ CH ₂ Cl
Vv	H	Me	CH ₂ CH ₂ Cl	H
Vw	OMe	Me	CH ₂ CH ₂ Cl	CH ₂ CH ₂ Cl
Vx	OMe	Me	CH ₂ CH ₂ Cl	H

The reaction is carried out at a temperature range of from 0°C to 100°C, preferably at room temperature, in a period from half an hour to eight hours, in a solvent such as N,N-dimethylformamide, N,N-dimethylacetamide diethyl ether or tetrahydrofuran, preferably in N,N-dimethylformamide.

In a similar manner, the corresponding 2-bromoethyl derivatives can be obtained by bromination of the 2-hydroxyethyl intermediates with thionyl bromide, phosphorus oxybromide or carbon tetrabromide and triphenylphosphine in N,N-dimethylformamide.

Acylation of the 2-hydroxyethyl intermediates with acid anhydride, such as acetic anhydride, benzoic anhydride or

methanesulfonic anhydride, or with acyl chloride, such as acetyl chloride, benzoyl chloride or methanesulfonyl chloride, in pyridine or in a mixture of chloroform and p-dimethylaminopyridine or methylenechloride and p-dimethylaminopyridine affords the corresponding acyl derivatives.

5

The following examples are illustrative of the process of the products of the present experiment, but are not to be construed as limiting.

10

EXAMPLE 1

To a hot solution of 1,8-dimethoxy-3-methyl-9,10-anthraquinone (4.5 g, 16 mM) and 1,3-dibromo-5,5-dimethylhydantoin (2.75 g, 19.2 mM) in carbon tetrachloride (500 mL) is added benzoyl peroxide (0.7 g), and the mixture is heated under reflux for 5 hours. The mixture is allowed to cool to room temperature. Insoluble hydantoin is removed by filtration, the filtrate is concentrated to dryness, and the residue recrystallized twice from ethyl acetate to give 3-bromomethyl-1,8-dimethoxy-9,10-anthraquinone (3.3 g, 57%), mp 176-178°C. ¹H NMR (CDCl₃) δ: 4.01 (3H, s, OMe), 4.03 (3H, s, OMe), 4.52 (2H, s, CH₂Br), 7.26-7.84 (5H, m, aromatic H). Analyses (C₁₇H₁₃BrO₄). Calculated: C, 56.53; H, 3.63; Br, 22.12. Found: C, 56.48; H, 3.67; Br, 21.93.

25

The mother liquors of recrystallization of above are concentrated, and the residue chromatographed on a silica gel column using a mixture of benzene and ethyl acetate. 3-Dibromomethyl-1,8-dimethoxy-9,10-anthraquinone (0.48 g) is eluted from the column followed by 3-bromomethyl-1,8-dimethoxy-9,10-anthraquinone (0.33 g). The former has the following characteristics: mp 207-210°C, ¹H NMR (CDCl₃) δ: 4.01 (3H, s, OMe), 4.07 (3H, s, OMe), 6.67 (1H, s, CHBr₂), 7.26-7.92 (5H, m, aromatic H). Analyses (C₁₇H₁₃Br₂O₄).

35

Calculated: C, 46.40; H, 2.75; Br, 36.31. Found: C, 46.63; H, 2.87; Br, 36.23.

Following the same procedure but using the corresponding anthraquinones as the starting materials, the following 3-bromomethyl- and 3,3-dibromomethyl-9,10-anthraquinones are

5 prepared:

- 3-Bromomethyl-1,6,8-trimethoxy-9,10-anthraquinone,
- 3-Dibromomethyl-1,6,8-trimethoxy-9,10-anthraquinone,
- 3-Bromomethyl-1,8-dihydroxy-9,10-anthraquinone,
- 3-Bromomethyl-1,8-dihydroxy-9,10-anthraquinone,
- 10 3-Bromomethyl-1,6,8-trihydroxy-9,10-anthraquinone,
- 3-Dibromomethyl-1,6,8-trihydroxy-9,10-anthraquinone,
- 3-Bromomethyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone,
- 3-Dibromomethyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone.

- 15 By the following the same procedure, but using N-bromosuccinimide instead of 3,3-dibromo-5,5-dimethylhydantoin, the same products above are prepared from their corresponding 3-methyl-9,10-anthraquinone starting materials.

20

EXAMPLE 2

- A mixture of 3-bromomethyl-1,8-dimethoxy-9,10-anthraquinone (1.05 g), 30% hydrogen bromide in acetic acid (5 mL) and
25 acetic acid (50 mL) is heated at 100°C for 5 hours. After cooling the mixture, 3-bromomethyl-1,8-dihydroxy-9,10-anthraquinone is collected by filtration, washed with acetic acid, and then air-dried to give 879 mg (91%) of the product, mp 220-222°C. ¹H NMR (CDCl₃) δ: 4.47 (2H, s, CH₂Br), 7.27-7.91 (5H, m, aromatic H), 12.02 (1H, s, OH), 12.05 (1H, s, OH). Analyses (C₁₅H₉BrO₄). Calculated: C, 54.08; H, 2.71; Br, 23.99. Found: C, 54.00; H, 2.92; Br, 24.16.

35

By following the same procedure but using the corresponding 1,8-dimethoxy-9,10-anthraquinones, the following compounds are prepared:

- 3-Dibromomethyl-1,8-dihydroxy-9,10-anthraquinone,
- 3-Bromomethyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone,
- 5 3-Dibromomethyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone.

EXAMPLE 3

A mixture of 3-bromomethyl-1,8-dihydroxy-9,10-anthraquinone
10 (456 mg) and bis(2-hydroxyethyl)amine (600 mg) in N,N-dimethylformamide (20 mL) is stirred for 2 hours, and then partitioned between chloroform (100 mL) and water (100 mL). The organic layer is separated, washed with water (50 mL x 3), dried over sodium sulfate, and then concentrated to
15 dryness. The residue is chromatographed on a silica gel column using chloroform-methanol (15:1 v/v) as the eluent. Upon concentration of the major fraction, 3-[N,N-bis(2-hydroxyethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone is obtained as a solid: ¹H NMR (CDCl₃) δ: 2.77 (4H, t,
20 NCH₂CH₂O), 3.78 (4H, t, NCH₂CH₂O), 3.78 (2H, s, CH₂N=), 7.21-7.79 (5H, m, aromatic H), 11.95 (1H, s, OH), 12.00 (1H, s, OH).

The solid is dissolved in 1N hydrochloric acid, and the
25 solvent is removed in vacuo. The crystalline hydrochloride salt (495 mg, 91%) is triturated with methanol (5 mL), mp 204-207°C (decomposition). Analyses (C₁₉H₁₉NO₆.HCL). Calculated: C, 57.95; H, 5.12; N, 3.56. Found: C, 58.00; H, 5.26; N, 3.37.

30

By following the same procedure but using the corresponding 3-bromomethyl-9,10-anthraquinones, the following compounds and their hydrochloric acid salts are prepared:

- 35 3-[N,N-Diethylamino]methyl-1,8-dihydroxy-9,10-anthraquinone,

- 3-[N-Ethylamino)methyl-1,8-dihydroxy-9,10-anthraquinone,
3-[N-(2-Hydroxyethyl)amino)methyl-1,8-dihydroxy-9,10-
anthraquinone,
3-(N,N-Diethylamino)methyl-1,8-dihydroxy-6-methoxy-
9,10-anthraquinone,
3-[N,N-Bis(2-hydroxyethyl)amino)methyl-1,8-dihydroxy
5 -6-methoxy-9,10-anthraquinone,
3-(N-Ethylamino)methyl-1,8-dihydroxy-6-methoxy-
9,10-anthraquinone,
3-[N-(2-Hydroxyethyl)amino)methyl-1,8-dihydroxy-6-methoxy-
9,10-anthraquinone,
10 3-(N,N-Diethylamino)methyl-1,8-dimethoxy-9,10-
anthraquinone,
3-[N,N-Bis(2-hydroxyethyl)amino)methyl-1,8-dimethoxy-
9,10-anthraquinone,
3-(N-Ethylamino)methyl-1,8-dimethoxy-9,10-anthraquinone,
15 3-[N-(2-Hydroxyethyl)amino)methyl-1,8-dimethoxy-9,10-
anthraquinone,
3-(N,N-Diethylamino)methyl-1,6,8-trimethoxy-9,10-
anthraquinone,
3-[N,N-Bis(2-hydroxyethyl)amino)methyl-1,6,8-trimethoxy-
20 9,10-anthraquinone,
3-(N-Ethylamino)methyl-1,6,8-trimethoxy-9,10-anthraquinone,
3-[N-(2-Hydroxyethyl)amino)methyl-1,6,8-trimethoxy-9,10-
anthraquinone,

25

EXAMPLE 4

To a solution of 3-[N,N-bis(2-hydroxyethyl)amino)methyl-1,8-
dihydroxy-9,10-anthraquinone, (162 mg) in dry N,N-
dimethylformamide (5 mL) is added thionyl chloride (0.2 mL).
30 After 2 hours at room temperature, the mixture is
concentrated in vacuo to dryness, and the residue is
trituated well with methanol (3 mL). 3-[N,N-Bis(2-
chloroethyl)amino)methyl-1,8-dihydroxy-9,10-anthraquinone
that is crystallized is collected by filtration, 172 mg
35 (96%), mp 211-214°C (decomposition). ¹H NMR (CDCl₃) δ: 3.48

(4H, t, $\text{NCH}_2\text{CH}_2\text{Cl}$), 3.85 (4H, t, $\text{NCH}_2\text{CH}_2\text{Cl}$), 4.34 (2H, d, $\text{CH}_2\text{N=}$), 7.32-7.95 (5h, M, aromatic H). Analyses ($\text{C}_{19}\text{H}_{17}\text{Cl}_2\text{NO}_4 \cdot \text{HCl}$). Calculated: C, 52.98; H, 4.21; Cl, 24.24; N, 3.25. Found: C, 52.79; H, 4.32; Cl, 24.41; N, 3.36.

By following the same procedure but using the corresponding (2-hydroxyethyl)amino derivatives, the following compounds
5 are prepared:

- 3-[N-(2-Chloroethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone,
3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dihydroxy-6
10 methoxy-9,10-anthraquinone,
3-[N-(2-Chloroethyl)amino]methyl-1,8-dihydroxy-6-methoxy-
9,10-anthraquinone,
3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dimethoxy-
9,10-anthraquinone,
15 3-[N-(2-Chloroethyl)amino]methyl-1,8-dimethoxy-
9,10-anthraquinone,
3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8,6-trimethoxy-
9,10-anthraquinone.
3-[N-(2-Chloroethyl)amino]methyl-1,6,8-trimethoxy-
20 9,10-anthraquinone,

Experimental Discussion

Table I list typical results supporting the use of the
25 present compounds as anti-cancer agents in the treatment of
subjects.

30

35

Table I. Inhibitory activity of some 9,10-anthraquinone derivatives.

	Compounds	melting point (°C)	ID ₅₀ (μg/mL)	ID ₅₀ (μM)
	IIa	194-195	>100	>390
	IIb	256-257	>100	>335
5	Va (HCl)	235-238(dec)	0.99	2.8
	Vb (HCl)	204-207(dec)	2.33	5.9
	Vc (HCl)	>275	0.26	0.77
	Vd (HCl)	255-261(dec)	0.066	0.16
	Ve (HBr)	240-241(dec)	0.51	1.16
	Vf (HCl)	225-227(dec)	5.80	13.7
	Vg (HBr)	>275	0.18	0.44
10	Vh (HCl)	259-260(dec)	0.072	0.19
	Vi (HCl)	154-158	52.3	128.3
	Vj (HCl)	202-205(dec)	>21	>49.8
	Vk (HCl)	254-255(dec)	1.30	3.59
	Vl (HCl)	251-252(dec)	53.7	142.0
	Vm (HCl)	222-223(dec)	2.90	6.91
	Vn (HCl)	226-227(dec)	12.5	27.7
15	Vo (HCl)	267-269(dec)	1.44	3.67
	Vp (HCl)	252-253(dec)	5.10	12.5
	Vq (HCl)	211-214(dec)	0.058	0.13
	Vr (HCl)	255-261(dec)	2.37	7.11
	Vs (HCl)	203-206(dec)	0.010	0.023
	Vu (HCl)	208-209(dec)	5.73	12.5
	Vw (HCl)	200-201(dec)	1.30	2.66

20

The starting materials, chrysophanol and emodin (IIa and IIb), are capable of intercalating into DNA but do not possess covalent bond forming capability, and exhibit little anticancer activity. The 1,8-dimethoxy intermediates (Vu-Vx), that bear alkylating potential but are incapable of intercalating into DNA due to the presence of bulky methoxy groups, are active to a small extent against mouse leukemia L1210 cells. Those compounds that are capable of intercalating into DNA and bind covalently to the DNA after intercalation (Vq-Vt) do exhibit extremely potent activity against L1210 cells.

30

Table II lists additional results supporting the anti-cancer use of the present experiment.

35

Table II. Antileukemic activity of chrysophanol derivatives bearing alkylating potential.

J and K		R ¹	R ²	R ³	L1210: ID ₅₀ , μM
5	Et	H	Me	Me	128.3
	Et	H	H(or Me)	Me (or H)	11.9
	Et	H	H	H	2.75
10	CH ₂ CH ₂ OH	H	Me	Me	>50
	CH ₂ CH ₂ OH	H	H(or Me)	Me (or H)	23.4
	CH ₂ CH ₂ OH	H	H	H	5.92
	CH ₂ CH ₂ Cl	H	Me	Me	12.5
	CH ₂ CH ₂ Cl	H	H(or Me)	Me (or H)	1.40
	CH ₂ CH ₂ Cl	H	H	H	0.13
15	Et	OMe	Me	Me	6.91
	Et	OMe	H	H	1.16
	CH ₂ CH ₂ OH	OMe	Me	Me	>27.6
20	CH ₂ CH ₂ OH	OMe	H	H	13.7
	CH ₂ CH ₂ Cl	OMe	Me	Me	2.66
	CH ₂ CH ₂ Cl	OMe	H(or Me)	Me (or H)	1.75
	CH ₂ CH ₂ Cl	OMe	H	H	0.023

25

3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone (Vq) is extremely potent against mouse leukemia L1210 made resistant to Cisplatin. At the dosage of 100 mg/kg/day x 5 (ip), mice inoculated with Cisplatin resistant leukemia L1210/Cisplatin are cured.

30

The process of treating tumors according to this invention comprises administering to a subject having an abnormal proportion of leukocytes or other evidence of a malignancy, a therapeutic nontoxic amount of a compound of the

35

experiment such as 3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone, as such or in the form of a pharmaceutically acceptable salt thereof. The invention also provides a pharmaceutical composition in dosage unit form comprising from 1 to 200 mg/kg of a compound of the invention, per dosage unit, together with pharmaceutically acceptable nontoxic inert carrier or diluent thereof as described above. A subject may be any warm-blooded animal and is preferably human.

EXPERIMENT 2:

Fifty-one new C-methyl-modified derivatives of the anthraquinones chrysophanol and emodin or their various methyl ethers were prepared for structure-activity relationship studies of anticancer activity against mouse leukemia L1210 and human leukemia HL-60 cells. Representative compounds were spectrophotometrically studied for their capacity to interact with natural and denatured DNA. In general, those anthraquinones bearing an amino function interact with DNA. 1,8-Dimethoxyanthraquinones are incapable of intercalating into DNA. 1- or 8-Monohydroxy-monomethoxyanthraquinones, however, interact with DNA to some extent. No straightforward correlation is apparent between the DNA-affinity data of the compounds studied spectrophotometrically and their cytotoxic effects. Cytotoxic potencies of these compounds on cell growth inhibition during a 72-h period are inversely correlated to their potencies when inhibiting [³H]TdR incorporation into DNA during the initial 30 min of exposure. Surprisingly, some compounds that showed more cytotoxicity did not inhibit initial TdR incorporation (0-30 min), while some others that strongly inhibited TdR incorporation initially did not exhibit cytotoxicity in 72 h. The results suggest that the cytotoxicity produced by these compounds is time dependent and is not a direct result of initial inhibition of DNA replication.

- It is well-known that one the of the metabolites of ellipticine, 9-hydroxyellipticine (Lesca, P., et al., Biochem. Pharmacol. 26:2169 (1977)) (9-OH-E), is also a potent anticancer agent. (Le Pecq, J.-B.; Cancer Res. 36:3067 (1976)) 2-N-Methyl-9-hydroxyellipticinium (9-OH-NME) is one of the most active drugs among the ellipticine analogues (Bernadou, J. Proc. Natl. Acad. Sci. U.S.A. 81:1297 (1984)). The latter is easily oxidized by peroxidases to 9-oxo-2-methylellipticinium (Auclair, C., et al., J. Med. Chem. 24:289 (1981); Bernadou, J., et al., J. Med. Chem. 26:574 (1983)) (9-oxo-NME), which is highly electrophilic and alkylates various nitrogen, (Meunier, G.; Tetrahedron Lett. 26:574 (1983); Auclair, C.; J. Med. Chem. 27:1161 (1984)) sulfur, (Monsarrat, B.; Biochem. Pharmacol. 32:3887 (1983)) and oxygen (Bernadou, J., et al., J. Med. Chem. 26:574 (1983); Meunier, G., et al., Tetrahedron Lett. 26:574 (1983)) nucleophiles. Among biological macromolecules, proteins, (Auclair, C., et al., Biochem. Pharmacol. 32:3883 (1984)) polyadenylate, (Dugue, B., et al., Biochem. Biophys. Res. Commun. 124:416 (1984)) RNA, (Dugue, B., et al.) and DNA (Auclair, C., et al., Biochemistry 25:1240 (1986)) are easily alkylated by 9-oxo-NME. This "biooxidative alkylation" has been proposed as a possible mode of anticancer action. (Dugue, B., et al., Cancer Res. 46:3828 (1986)).
- For other intercalating anticancer agents, such as amsacrine (m-AMSA), and the anthracycline antibiotics, extensive studies on their mechanism of anticancer action (Nelson, E.M., et al., Proc. Natl. Acad. Sci. U.S.A. 81:1361 (1984); Chen, G.L., et al., J. Biol. Chem. 259:13560 (1984); Riou, J.F., et al., Biochem. Pharmacol. 35:4409 (1986); Crooke, S.T., Reich, S.D.; Eds Anthracyclines: Current Status and New Developments; Academic Press: New York, (1980); Myers, C.E. In Pharmacologic Principles of Cancer Treatment; Chabner, B., Ed.; Saunders: Philadelphia p. 416 (1982); DiMarco, A. In Cancer Medicine; Holland, J.F., Frei, E. Eds;

Lea and Febiger: Philadelphia, 2nd ed., p. 872 (1982); Muggia, F.M., Young, C.W., Carter, S.K. Eds. Anthracycline Antibiotics in Cancer Therapy; Martinus Nijhoff: Hague/Boston/New York, pp. 71-174 (1982)) and QSAR studies directed toward the development of more selective drugs have been conducted. (Ferguson, L.R., Denny, W.A., J. Med. Chem. 5 23:269 (1980); Denny, W.A., et al., J. Med. Chem. 25:276 (1982); Alexander, J., et al., J. Med. Chem. 27:1343 (1984)). Whether these intercalators bind covalently to biomolecules has not been established.

10 In order to test our hypothesis that intercalating agents with covalent bond forming capability may exert potent cytocidal activity, we chose chrysophanol Ia and emodin Ib (Figure 1) as the starting materials; Ia and Ib were isolated from crude rhubarb extract. (Kelly, T.R.; J. Org. 15 Chem. 48:3573 (1983)). Both compounds exhibit little anticancer or cytocidal activity. Their structural aromatic features indicate that these compounds and their derivatives (particularly positively charged ones) may intercalate into double helix of nucleic acids.

20

Chemistry

The hydroxy groups at the 1- and 8-positions of I were methylated with methyl sulfate and K_2CO_3 in acetone 25 (Alexander, J., et al., J. Org. Chem. 45:20 (1980)) to the known 1,8-dimethoxy-9,10-anthraquinones II. The C-methyl group of II was then brominated with NBS (Banvill, J., et al., J. Chem Soc. Perkin Trans. I, 613 (1976)) or 1,3-dibromo-5,5-dimethylhydantoin (BDH) (Sargent, M.V., et al., 30 J. Chem. Soc., C 2763 (1969); Cava, M.P., et al., Tetrahedron 40:4767 (1984)) in carbon tetrachloride in the presence of benzoyl peroxide to give monobromide III as the major product along with a small amount of dibromide IV. Treatment of III with various amines including mono(2- 35 hydroxyethyl)amine and bis(2-hydroxyethyl)amine afforded the

corresponding alkyl-amino derivatives IX (Table III): Chlorination of the (2-hydroxyethyl)amino derivatives IX-2 gave the corresponding 3-[[[(2-chloroethyl)amino]methyl]-9,10-anthraquinones IX-3. On the basis of reports by Anderson et al. (Anderson, W.K., et al., J. Med. Chem. 20:812 (1977); Anderson, W.K., et al., J. Med. Chem. 22:977 (1979); Anderson, W.K., et al., J. Med. Chem. 26:1333 (1983); Zwelling, L.A. Cancer Metastasis Rev. 4:263 (1985)) that certain carbamates are susceptible to nucleophilic attack, we synthesized N-methylcarbamate IX-4 by treatment of IX-2 with N-methyl isocyanate.

10

15

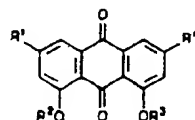
20

25

30

35

Table III
Synthetic Derivatives of Chrysophanol and Emodin



compd ^a	R ¹	R ²	R ³	R ⁴	mp, °C ^b	formula
5						
Ia	H	Me	Me	Me	189-190	C ₂₇ H ₃₀ O ₄
Ib	OMe	Me	Me	Me	228-229	C ₂₈ H ₃₂ O ₅
IIa	H	Me	Me	CH ₂ Br	176-178	C ₂₇ H ₂₈ BrO ₄
IIb	OMe	Me	Me	CH ₂ Br	250-254	C ₂₈ H ₃₀ BrO ₅
IVa	H	Me	Me	CHBr ₂	207-210	C ₂₇ H ₂₆ Br ₂ O ₄
IVb	OMe	Me	Me	CHBr ₂	254-257	C ₂₈ H ₂₈ Br ₂ O ₅
Va	H	H or Me	Me or H	CH ₂ Br	213-215	C ₂₇ H ₂₈ BrO ₄
Vb	OMe	H or Me	Me or H	CH ₂ Br	199-201	C ₂₈ H ₃₀ BrO ₅
VIa	H	H	H	CH ₂ Br	220-222	C ₂₇ H ₂₈ BrO ₄
VIb	OMe	H	H	CH ₂ Br	248-250	C ₂₈ H ₃₀ BrO ₅
VIIa	H	H or Me	Me or H	CHBr ₂	176-178	C ₂₇ H ₂₆ Br ₂ O ₄
VIIb	OMe	H or Me	Me or H	CHBr ₂	202-203	C ₂₈ H ₂₈ Br ₂ O ₅
VIIIa	H	H	H	CHBr ₂	211-213	C ₂₇ H ₂₆ Br ₂ O ₄
VIIIb	OMe	H	H	CHBr ₂	237-238	C ₂₈ H ₂₈ Br ₂ O ₅
10						
IXa-1	H	Me	Me	CH ₂ NEt ₂	154-158	C ₂₇ H ₃₀ NO ₄ ·HCl·H ₂ O
IXa-2	H	Me	Me	CH ₂ N(CH ₂ CH ₂ OH) ₂	202-205d	C ₂₇ H ₃₂ NO ₆ ·HCl
IXa-3	H	Me	Me	CH ₂ N(CH ₂ CH ₂ Cl) ₂	205-206d	C ₂₇ H ₂₈ Cl ₂ NO ₄ ·HCl
IXa-4	H	Me	Me	CH ₂ N(CH ₂ CH ₂ OCONHMe) ₂	120d	C ₂₇ H ₃₂ N ₂ O ₇ ·HCl
IXa-5	H	Me	Me	CH ₂ NHEt	254-255d	C ₂₇ H ₃₀ NO ₄ ·HCl·H ₂ O
IXa-6	H	Me	Me	CH ₂ NHCH ₂ CH ₂ OH	251-252d	C ₂₇ H ₃₂ NO ₆ ·HCl
IXa-7	H	Me	Me	CH ₂ NHCH ₂ CH ₂ Cl	208-209d	C ₂₇ H ₂₈ ClNO ₄
IXb-1	OMe	Me	Me	CH ₂ NEt ₂	222-223d	C ₂₈ H ₃₂ NO ₅ ·HCl
IXb-2	OMe	Me	Me	CH ₂ N(CH ₂ CH ₂ OH) ₂	225-227d	C ₂₈ H ₃₄ NO ₇ ·HCl
IXb-3	OMe	Me	Me	CH ₂ N(CH ₂ CH ₂ Cl) ₂	200-201d	C ₂₈ H ₃₀ Cl ₂ NO ₅ ·HCl
IXb-5	OMe	Me	Me	CH ₂ NHEt	267-269d	C ₂₈ H ₃₀ NO ₅ ·HCl
IXb-6	OMe	Me	Me	CH ₂ NHCH ₂ CH ₂ OH	252-253d	C ₂₈ H ₃₂ NO ₇ ·HCl
IXb-7	OMe	Me	Me	CH ₂ NHCH ₂ CH ₂ Cl·HCl·DMF	204-205d	C ₂₈ H ₃₀ ClNO ₅ ·HCl·C ₂ H ₅ NO
Xa-1	H	H or Me	Me or H	CH ₂ NEt ₂	225-227d	C ₂₇ H ₃₀ NO ₄ ·HBr
Xa-2	H	H or Me	Me or H	CH ₂ N(CH ₂ CH ₂ OH) ₂	209-216d	C ₂₇ H ₃₂ NO ₆ ·HCl
Xa-3	H	H or Me	Me or H	CH ₂ N(CH ₂ CH ₂ Cl) ₂	203-205d	C ₂₇ H ₂₈ Cl ₂ NO ₄ ·HCl
Xa-4	H	H or Me	Me or H	CH ₂ N(CH ₂ CH ₂ OCONHMe) ₂	178-182d	C ₂₇ H ₃₂ N ₂ O ₇ ·HCl
Xb-1	OMe	H or Me	Me or H	CH ₂ NEt ₂	110-112	C ₂₈ H ₃₂ NO ₅
Xb-2	OMe	H or Me	Me or H	CH ₂ N(CH ₂ CH ₂ OH) ₂	221-223	C ₂₈ H ₃₄ NO ₇ ·HCl
Xb-3	OMe	H or Me	Me or H	CH ₂ N(CH ₂ CH ₂ Cl) ₂	152-154	C ₂₈ H ₃₀ Cl ₂ NO ₅
20						
XIa-1	H	H	H	CH ₂ NEt ₂	236-238d	C ₂₇ H ₃₀ NO ₄ ·HCl·1/2H ₂ O
XIa-2	H	H	H	CH ₂ N(CH ₂ CH ₂ OH) ₂	204-207d	C ₂₇ H ₃₂ NO ₆ ·HCl
XIa-3	H	H	H	CH ₂ N(CH ₂ CH ₂ Cl) ₂	211-214d	C ₂₇ H ₂₈ Cl ₂ NO ₄ ·HCl
XIa-4	H	H	H	CH ₂ N(CH ₂ CH ₂ OCONHMe) ₂	125-131	C ₂₇ H ₃₂ N ₂ O ₇ ·HCl
XIa-5	H	H	H	CH ₂ NHEt	> 275	C ₂₇ H ₃₀ NO ₄ ·HCl
XIa-6	H	H	H	CH ₂ NHCH ₂ CH ₂ OH	255-261d	C ₂₇ H ₃₂ NO ₆ ·HCl
XIa-7	H	H	H	CH ₂ NHCH ₂ CH ₂ Cl	255-261d	C ₂₇ H ₂₈ ClNO ₄ ·HCl
XIa-8	H	H	H	CH ₂ NH ₂	240-245d	C ₂₇ H ₂₈ NO ₄ ·HCl·H ₂ O
XIa-9	H	H	H	CH ₂ NMe ₂	282-283d	C ₂₇ H ₃₀ NO ₄ ·HCl
XIa-10	H	H	H	CH ₂ NH(CH ₂) ₂ Me	> 275	C ₂₇ H ₃₂ NO ₅ ·HCl
XIa-11	H	H	H	CH ₂ NHCH ₂ CH ₂ CH ₂ OH	251-252d	C ₂₇ H ₃₄ NO ₇ ·HCl
XIa-12	H	H	H	CH ₂ N(CH ₂ CH ₂ OCONH·iPr) ₂	159-160	C ₂₇ H ₃₈ N ₂ O ₇ ·HCl
XIa-13	H	H	H	CH ₂ N(CH ₂) ₃	255-257d	C ₂₇ H ₃₀ NO ₄ ·HCl
XIa-14	H	H	H	CH ₂ N(CH ₂) ₃	246-247d	C ₂₇ H ₃₀ NO ₄ ·HCl
XIa-18	H	H	H	CH ₂ (imidazol-1-yl)	270-272d	C ₂₇ H ₂₈ N ₂ O ₄ ·HCl·H ₂ O
25						
XIb-1	OMe	H	H	CH ₂ NEt ₂	240-241d	C ₂₈ H ₃₂ NO ₅ ·HBr
XIb-2	OMe	H	H	CH ₂ N(CH ₂ CH ₂ OH) ₂	225-227d	C ₂₈ H ₃₄ NO ₇ ·HCl
XIb-3	OMe	H	H	CH ₂ N(CH ₂ CH ₂ Cl) ₂	203-206d	C ₂₈ H ₃₀ Cl ₂ NO ₅ ·HCl
XIb-5	OMe	H	H	CH ₂ NHEt	> 275	C ₂₈ H ₃₀ NO ₅ ·HBr
XIb-6	OMe	H	H	CH ₂ NHCH ₂ CH ₂ OH	259-260d	C ₂₈ H ₃₂ NO ₇ ·HCl
XIb-7	OMe	H	H	CH ₂ NHCH ₂ CH ₂ Cl	powder	C ₂₈ H ₃₀ ClNO ₅

^a All the compounds were analyzed for C, H, X (Br or Cl), and/or N. Analyses for these elements were within ±0.4% of the theoretical values required unless specified otherwise. ^b For nitrogen-containing compounds, melting points were of the HX salt. ^d = decomposition.

^c Unstable, and satisfactory analyses could not be obtained.

The methyl protecting groups at 1 and 8 could be removed stepwise at various stages (Figure 1). Thus, treatment of IX with HBr in acetic acid at room temperature afforded a crystalline mixture of 1-O-methyl- and 8-O-methyl-anthraquinones X, whereas acid hydrolysis at reflux temperature for a few hours resulted in complete demethylation, giving rise to XI. Later, it was found that partially methylated chrysophanol and emodin could be directly brominated to the corresponding mixtures of the monobromides V (major products) and dibromides VII. The former were treated with amines to give X, which were further converted to the corresponding XI. Alternatively, the 3-[(alkylamino)methyl] derivatives XI were prepared by amination of VI. Chlorination of XIa-2 with thionyl chloride afforded XIa-3. N-Methyl- and N-isopropyl-carbamates, XIa-4 and XIa-12, respectively, were prepared by treatment of XIa-2 with the corresponding N-alkyl isocyanates.

Spectrophotometric Studies for DNA Interactions

Some representative compounds were studied for their ability to interact with nucleic acids in solution by comparison of the electronic spectrum of drug alone with that of the drug in the presence of an excess of nucleic acid. All drugs studied have an absorption band in the visible region, separate from the absorption band of the nucleic acids, and therefore any changes in band intensity and position were indicative of drug chromophore-DNA interaction. It was observed that both chrysophanol (Ia) and emodin (Ib) and their derivatives lacking the basic center do not appear to interact with DNA to any significant extent. Since these compounds have low solubility in aqueous solutions, the spectral measurement alone may not be sufficient however to allow one to draw a definite conclusion.

Generally, those anthraquinones bearing an amino function (e.g., Xa-1, Xb-1, XIa-1, XIa-2, XIb-2, and XIb-6) interact with both native and thermally denatured DNA but more strongly with native DNA (Figure 2). As expected, 1-8-di-O-methylchrysophanol and 1,6,8-tri-O-methylemodin analogues IXa and IXb do not interact with DNA. Anthraquinones wherein one peri-hydroxyl group is methylated (e.g., Xa-2 and Xb-2) interact with DNA to a lesser extent than the corresponding unmethylated (XIa-2 and XIb-2). It is interesting to note that while DNA induced changes in absorption spectra of some derivatives (e.g., XIa-7 and XIb-7), these do not appear to be connected with the alkylating capabilities in view of the fact that they could be reversed by addition of Me₂SO (to 1:1 v/v). The dissociation of nonbonded ligand-DNA complexes in the presence of organic solvents is a phenomenon well documented. (Wakelin, L.P.G., et al., *Biochemistry* 20:5779 (1981)).

Changes in the absorption spectra of the drugs (Figure 2) are not inconsistent with the possibility of the intercalative mode of binding. Other types of nonbinding interactions, however, (e.g. binding to the minor groove of the double helix (Bloomfield, V.A., et al., *In Physical Chemistry of Nucleic Acids*; Harper and Row: New York, pp. 432-434 (1974); Jorgenson, K.F., et al., *J. Biomol. Struct. Dyn.* 6:1005 (1988)) cannot be excluded. It is well-known that intercalative binding, which most often has an ionic component, is affected by a rise in concentration of salts, (Kapuscinski, J., Darzynkiewicz, Z.J., *Biomol. Struct. Dyn.* 5:127 (1987), and compounds such as Xa-2, Xb-2, XIb-3, and XIb-7 lost the ability to interact with DNA when Na⁺ concentration was increased from 0.01 to 0.1 M. On the basis of this fact one can conclude that the affinity for DNA of these compounds is lower than those that interact with DNA at both ionic strengths.

No straightforward correlation is apparent between the DNA-affinity data of the drugs studied and their biological activity.

Biological Activities

5 Preliminary biological data for inhibiting cell growth of murine L1210 leukemic cells and human acute promyelocytic leukemia cells (HL-60) during 72 h of exposure to the compounds are given in Table IV. The potencies for inhibiting [³H]TdR incorporation into DNA in HL-60 cells during the initial 30-min period are also given in Table IV. 10 It is interesting to note that 1,8-di-O-methyl derivatives are uniformly devoid of activity against L1210 leukemic cells. These results are consistent with published data that compare the ratios of the potencies (ID₅₀'s) for cell growth inhibition (A) and inhibition of [³H]TdR 15 incorporation into DNA in HL-60 cells (B). The B/A ratios allow an indirect estimation of whether or not cytotoxicities exerted by these analogues are primarily due to initial inhibition of DNA synthesis. The B/A ratios for Vb, VIIa, VIIb, VIIIb, Xa-3, and XIa-3 are 41, 217, 237, 20 579, 39, and 383, respectively, suggesting that these compounds exert their initial effects mainly on processes other than DNA synthesis per se (Table V). These results suggest that these compounds exert their cytotoxic effects in a time-dependent manner and their initial action is 25 targeted at the sites other than DNA elongation. Whether these analogues, like m-AMSA, ellipticine, or anthracyclines, act by inhibiting DNA topoisomerase II remains to be explored. It is of interest to note that the above-mentioned anthraquinones are among the most potent 30 antileukemic analogues listed in Table IV, with ID₅₀ values ranging from 1.4×10^{-6} to 4.2×10^{-11} for L1210⁴⁸ cells and 1.7×10^{-6} to 6.0×10^{-8} M for HL-60 cells. Our preliminary experiments indicate that the compounds arrest cells in the S and/or G₂ phases of the cell cycle (unpublished results). 35

TABLE IV
Biological Activities of Derivatives of
Chrysophanol and Emodin

		ID ₅₀ M (L1210 cell growth)	ID ₅₀ M (HL-60 cell growth (72 h)) (A)	ID ₅₀ M (HL-60 TdR into DNA) (B)	B/A
5	IIa	2.8×10^{-4}	1.0×10^{-4}	1.9×10^{-4}	1.9
	IIb	1.0×10^{-4}	4.9×10^{-4}	2.1×10^{-4}	0.43
	IIIa	9.2×10^{-4}	4.4×10^{-4}	1.1×10^{-4}	2.5
	IIIb	6.8×10^{-7}	8.4×10^{-4}	1.2×10^{-4}	0.14
	IVa	8.9×10^{-7}	2.1×10^{-4}	9.0×10^{-4}	4.3
	IVb	4.1×10^{-4}	6.0×10^{-4}	5.8×10^{-4}	9.7
	Va	8.8×10^{-4}	7.1×10^{-4}	1.2×10^{-4}	1.7
	Vb	6.8×10^{-4}	1.7×10^{-4}	7.0×10^{-4}	41
	VIa	8.6×10^{-4}	7.1×10^{-4}	1.6×10^{-4}	2.3
	VIb	5.9×10^{-4}	2.5×10^{-4}	3.4×10^{-4}	13.6
10	VIIa	4.2×10^{-11}	6.0×10^{-4}	1.3×10^{-4}	217
	VIIb	1.0×10^{-9}	9.7×10^{-4}	2.3×10^{-4}	237
	VIIIa	4.4×10^{-7}	2.5×10^{-4}	2.9×10^{-4}	18.6
	VIIIb	1.0×10^{-9}	1.9×10^{-7}	1.1×10^{-4}	579
	IXa-1	1.3×10^{-4}	3.0×10^{-4}	1.9×10^{-4}	6.3
	IXa-2	$>5.0 \times 10^{-4}$	4.9×10^{-4}	1.4×10^{-4}	0.03
	IXa-3	1.3×10^{-4}	1.8×10^{-4}	1.4×10^{-4}	7.8
	IXa-4	7.2×10^{-4}	9.5×10^{-4}	9.4×10^{-4}	0.99
	IXa-5	1.2×10^{-4}	4.8×10^{-4}	2.9×10^{-4}	0.60
	IXa-6	1.4×10^{-4}	5.5×10^{-4}	4.6×10^{-4}	0.84
15	IXa-7	2.6×10^{-4}	8.6×10^{-4}	1.9×10^{-4}	0.22
	IXb-1	6.9×10^{-4}	7.9×10^{-4}	4.2×10^{-4}	0.53
	IXb-2	$>2.7 \times 10^{-4}$	1.0×10^{-4}	1.7×10^{-4}	0.17
	IXb-3	2.7×10^{-4}	2.0×10^{-4}	8.9×10^{-4}	4.5
	IXb-5	9.8×10^{-4}	1.7×10^{-4}	1.3×10^{-4}	0.76
	IXb-6	1.7×10^{-4}	1.1×10^{-4}	2.6×10^{-4}	0.24
	IXb-7	3.2×10^{-4}	5.8×10^{-4}	1.8×10^{-4}	3.1
	Xa-1	1.2×10^{-4}	5.2×10^{-4}	1.4×10^{-4}	2.7
	Xa-2	$>2.4 \times 10^{-4}$	2.1×10^{-4}	8.9×10^{-4}	4.2
	Xa-3	1.4×10^{-4}	3.9×10^{-7}	1.5×10^{-4}	38.5
20	Xa-4	8.9×10^{-4}	1.2×10^{-4}	1.2×10^{-4}	1
	Xb-1	6.9×10^{-4}	6.7×10^{-4}	1.8×10^{-4}	26.9
	Xb-2	8.8×10^{-4}	$>5.0 \times 10^{-4}$	9.6×10^{-4}	0.02
	Xb-3	3.3×10^{-4}	5.2×10^{-7}	5.4×10^{-4}	10.4
	XIa-1	2.8×10^{-4}	1.8×10^{-4}	1.4×10^{-4}	7.8
	XIa-2	5.9×10^{-4}	3.3×10^{-4}	1.4×10^{-4}	4.2
	XIa-3	1.3×10^{-7}	1.8×10^{-7}	6.9×10^{-4}	383
	XIa-4	$>1.8 \times 10^{-4}$	2.1×10^{-4}	2.8×10^{-4}	0.13
	XIa-5	7.7×10^{-7}	8.7×10^{-4}	1.8×10^{-4}	2.1
	XIa-6	1.6×10^{-7}	8.6×10^{-7}	1.5×10^{-4}	17.4
25	XIa-7	7.1×10^{-4}	7.5×10^{-4}	3.9×10^{-4}	5.2
	XIa-8	8.5×10^{-4}	1.1×10^{-4}	2.5×10^{-4}	2.3
	XIa-9	2.2×10^{-4}	2.7×10^{-4}	9.3×10^{-4}	3.3
	XIa-10	4.6×10^{-4}	4.1×10^{-4}	2.5×10^{-4}	6.1
	XIa-11	6.7×10^{-7}	3.2×10^{-4}	1.6×10^{-4}	5.0
	XIa-12	2.7×10^{-4}	2.7×10^{-4}	1.7×10^{-4}	0.63
	XIa-13	1.8×10^{-4}	2.1×10^{-4}	1.0×10^{-4}	4.8
	XIa-14	4.0×10^{-4}	2.8×10^{-4}	6.3×10^{-4}	22.5
	XIa-15	7.4×10^{-4}	4.8×10^{-4}	4.3×10^{-4}	9.0
	XIb-1	1.2×10^{-4}	3.4×10^{-4}	3.5×10^{-4}	10.3
30	XIb-2	1.4×10^{-4}	6.6×10^{-4}	4.5×10^{-4}	5.2
	XIb-3	2.3×10^{-4}	6.1×10^{-7}	1.2×10^{-4}	19.7
	XIb-5	7.2×10^{-7}	2.1×10^{-4}	1.3×10^{-4}	6.2
	XIb-6	5.0×10^{-7}	1.6×10^{-4}	1.2×10^{-4}	7.5
	XIb-7	1.8×10^{-4}	9.2×10^{-4}	2.2×10^{-4}	23.9

TABLE V

Inverse Relationship Between Cell Growth Inhibition
and Inhibition of Initial Thymidine Incorporation Into
DNA in HL-60 Cells by Chrysophanol Derivatives^a

no. of compds examined	value of IC ₅₀ (cell growth), μ M		ratio of IC ₅₀ (dThd incorpn)/ IC ₅₀ (cell growth), mean \pm SE
	range	mean \pm SE	
8	<1	0.36 \pm 0.10	167.3 \pm 72.4
17	1-5	2.73 \pm 0.24	9.55 \pm 2.30
15	5-10	7.42 \pm 0.53	6.32 \pm 2.10
7	10-50	28.43 \pm 5.51	3.03 \pm 1.83
8	>50	204.4 \pm 65.4	0.22 \pm 0.09

^aCell growth inhibition was measured at the end of 72-h exposure to each compound as described under Experimental Section. Inhibition of [³H]dThd incorporation into DNA was measured during the first 30 min of exposure to each corresponding compound as described under Experimental Section.

Materials and Methods

Melting points were determined on a Thomas-Hoover capillary apparatus and are uncorrected. Elemental analyses were performed by M-H-W Laboratories, Phoenix, AZ, and all new compounds with the exception of IXa-7 and XIa-7, which were unstable, analyzed correctly. ¹H NMR spectra were recorded on a JEOL FX90Q spectrometer with Me₄Si as the internal standard. Chrysophanol (Ia) and emodin (Ib) were isolated from rhubarb extract by the procedure of Kelly et al. (Kelly, T.R., et al. J. Org. Chem. 48:3573 (1983)) except CH₂Cl₂ was used instead of Et₂O throughout the isolation process.

The following examples are illustrative of the process and products of the present invention, but are not to be construed as limiting.

EXAMPLE 5

5 **1,8-Dimethoxy-3-methyl-9,10-anthraquinone** (1,8-Di-O-methylchrysophanol, IIa). A mixture of chrysophanol (Ia, 7.0 g, 0.029 mol), K_2CO_3 (10 g, 0.071 mol), and Me_2SO_4 (10 mL, 0.1 mol) in Me_2CO (300 mL) was stirred under reflux for 16 h and then concentrated in vacuo. The residue was trituated well with water (300 mL), and the crystalline IIa (7.5 g, 96%) was collected by filtration and air-dried: mp 191-193 °C (lit. (Beilstein, 8:473) mp 195 °C); 1H NMR ($CDCl_3$) δ 2.46 (3 H, s, 3-Me), 3.98 (3 H, s, OMe), 3.99 (3 H, s, OMe), 7.08-7.86 (5 H, m, H-2,4,5,6,7). Anal. ($C_{17}H_{14}O_4$) C, H.

EXAMPLE 6

15 **1,3,8-Trimethoxy-6-methyl-9,10-anthraquinone** (1,3,8-Tri-O-methylemodin, IIb). In a similar manner, emodin (Ib, 1.0 g, 3.9 mmol) was converted into IIb (1.04 g, 90%): mp 225 °C (lit. (Beilstein, 8:523) mp 225 °C); 1H NMR (Me_2SO-d_6) δ 2.43 (3 H, s, 6-Me) 3.88 (3 H, s, OMe), 3.92 (6 H, s, 2 x OMe), 20 6.94 (1 H, d, H-7, $J_{5,7} = 2.2$ Hz), 7.13 (1 H, d, H-5, $J_{5,7} = 2.2$ Hz), 7.33 (1 H, s, H-2), 7.46 (1 H, s, H-4). Anal. ($C_{18}H_{16}BrO_5$) C, H.

EXAMPLE 7

25 **3-(Bromomethyl)-1,8-dimethoxy-9,10-anthraquinone (IIIa)**. To a hot solution of IIa (4.5 g, 0.016 mol) and BDH (2.75 g, 0.019 mol) in CCl_4 (500 mL) was added Bz_2O_2 (0.7 g), and the mixture was heated under reflux for 5 h. The mixture was allowed to cool room temperature. Insoluble hydantoin was removed by filtration, the filtrate was concentrated in vacuo, and the residue was crystallized twice from EtOA c to give IIIa (3.3 g, 57%): mp 176-178 °C (lit. (Blankespoor, R.L., et al., J. Org. Chem. 52:2059 (1987) mp 174-175 °C); 1H NMR ($CDCl_3$) δ 4.01 (3 H, s, OMe), 4.03 (3 H, s, OMe), 4.01 30 35

(3 H, s, OMe), 4.52 (2 H, s, CH_2Br), 7.26-7.84 (5 H, m, H-2,4,5,6,7). Anal. ($\text{C}_{17}\text{H}_{13}\text{BrO}_4$) C, H, Br.

EXAMPLE 8

3-(Dibromomethyl)-1,8-dimethoxy-9,10-anthraquinone (IVa). The mother liquors of recrystallization were concentrated, and the residue was chromatographed on a silica gel column using a mixture of C_6H_6 and EtOAc (3:1) to give IVa (0.48 g): mp 207-210 degrees C; ^1H NMR (CDCl_3) δ 4.01 (3 H, s, OMe), 4.07 (3 H, s, OMe), 6.67 (1 H, s, CHBr_2), 7.26 (5 H, m, H-2,4,5,6,7). Anal ($\text{C}_{17}\text{H}_{13}\text{Br}_2\text{O}_4$) C, H, Br.

A further amount of IIIa (0.33 g) was eluted from the column, making the total yield of 62.7%.

EXAMPLE 9

6-(Dibromomethyl)-1,3,8-trimethoxy-9,10-anthraquinone (IVb). In a similar manner, IIb (3.12 g, 0.01 mol) was brominated to give IIIB (3.06 g, 74.4%) [mp 250-254 °C; (lit. Alexander, J., et al., J. Org. Chem. 45:20 (1980)) mp. 233.5-234 degrees C]; ^1H NMR (CDCl_3) was identical with that reported (Alexander, J., et al.) and IVb (197 mg) [mp 254-257 degrees C; ^1H NMR (CDCl_3) δ 3.95 (6 H, s, 2 X OMe), 3.97 (3 H, s, OMe), 6.67 (1 H, s, CHBr_2), 6.78 (1 H, d, H-7, $J_{5,7} = 2.47$ Hz), 7.32 (1 H, d, H-5), 7.54 (1 H, d, H-2, $J_{2,4} = 1.92$ Hz), 7.90 (1 H, d, H-4)]. Anal. ($\text{C}_{17}\text{H}_{14}\text{Br}_2\text{O}_5$) C, H, Br.

EXAMPLE 10

3-(Bromomethyl)-1(and 8)-Hydroxy-8(and 1)-Methoxy-9,10-anthraquinone (Va). A mixture of IIIa (70 mg, 0.19 mmol in HOAc (10 mL) and 30% HBr/HOAc (1 mL) was stirred overnight at room temperature and then concentrated in vacuo. The residue was chromatographed on a silica gel column using CHCl_3 as the eluent to give 51 mg (76%) of Va as yellow crystals: mp 213-215 °C; ^1H NMR (CDCl_3) δ 4.08, 4.10 (2 x 3 H, 2 s, 1- and 8-OMe), 4.47, 4.54 (2 X 2 h, 2 s, CH_2Br), 7.24-8.04 (10 H, m, H-2,4,5,6,7). Anal. ($\text{C}_{16}\text{H}_{11}\text{BrO}_4$) C, H, Br.

EXAMPLE 11

3-(Dibromomethyl)-1(and 8)-hydroxy-8(and 1)-methoxy-9,10-anthraquinone (VIIa) and 6-(Dibromomethyl)-1(and 8)-hydroxy-3,8(and 1,3)-dimethoxy-9,10-anthraquinone (VIIf). In a similar manner, from IVa (200 mg, 0.453 mmol) and IVb (100 mg, 0.213 mmol), VIIa (82 mg, 42.5%) and VIIf (82 mg., 84.4%), respectively, were prepared (see Table III).

5

EXAMPLE 12

3-(Bromomethyl)-1,8-dihydroxy-9,10-anthraquinone (VIa). A mixture of III a (1.05 g, 2.9 mmol), HOAc (50 mL), and 30% HBr/HOAc (5 mL) was heated at 100 °C for 5 h. After the mixture was cooled, VIa was collected by filtration, washed with HOAc and H₂O, and then air-dried to give 879 mg (91%) of the product: mp 220-222 °C; ¹H NMR (CDCl₃) δ 4.47 (2 H, s, CH₂Br), 7.22-7.90 (5 H, m, H-2,4,5,6,7), 12.01, 12.04 (2 x H, 2 s, 1-OH, 8-OH). Anal. (C₁₅H₉BrO₄) C, H, Br.

10

15

EXAMPLE 13

3-(Dibromomethyl)-1,8-dihydroxy-9,10-anthraquinone (VIIIa) and 6-(Dibromomethyl)-3-methoxy-1,8-dihydroxy-9,10-anthraquinone (VIIf). In a similar manner, IVa (237 mg, 0.54 mmol) and IVb (472 mg, 1 mmol) were converted into VIIIa (166 mg, 75%) and VIIf (408 mg, 95%), respectively (Table III).

20

EXAMPLE 14

3-[(Diethylamino)methyl]-1,8-dimethoxy-9,10-anthraquinone (IXa-1). To a solution of IIIa (200 mg, 0.55 mmol) in DMF (10.0 mL) was added Et₂NH (5.0 mL), and the mixture was stirred at room temperature for 3 days. The mixture was partitioned between EtOAc (20 mL) and H₂O (20 mL). The aqueous layer was washed with EtOAc (20 mL). The combined EtOAc solutions were washed with H₂O (2 x 20 mL) and saturated NaCl (2 x 20 mL), dried (Na₂SO₄), and concentrated, and the residue was chromatographed on a silica gel column first with CHCl₃, which eluted the 3-(hydroxymethyl)

25

30

35

derivative (33 mg), followed by CHCl_3 containing 3% MeOH. The IXa (140 mg) that eluted from the column was converted to the crystalline HCl salt (142 mg, 63%), mp 154-158 °C.

In a similar manner but by using the corresponding amines, IXa-2, -5, and -6, were prepared (Table III). Also by use of the same procedure starting from IIIb and the corresponding amines, IXb-1, -2, -5, and -7 were synthesized (Table III).

EXAMPLE 15

3-[(Diethylamino)methyl]-(or 8)-hydroxy-8(or 1)-methoxy-9,10-anthraquinone (Xa-1). The HCl salt monohydrate of IXa (100 mg, 0.25 mmol) was dissolved in a mixture of HOAc (5 mL) and 30% HBr/HOAc (0.4 mL), and the solution was stirred at room temperature for 24 h. After concentration in vacuo, the residue was partitioned between saturated NaHCO_3 (10 mL) and CHCl_3 (10 mL). The CHCl_3 layer was dried (Na_2SO_4) and concentrated and the residue chromatographed on a silica gel column using CHCl_3 -MeOH (30:1 v/v) to give Xa-1 as a glass, which was dissolved in 1 N HBr (1 mL). Upon dilution of the solution with EtOH (5 mL), the monohydrobromide of Xa-1 (82 mg) precipitated as yellow microcrystals, mp 225-227 °C dec.

In a similar manner, Xa-2-4 and Xb 1-4 were prepared (Table III).

EXAMPLE 16

3-[[N,N-Bis(2-hydroxyethyl)amino]methyl]-1,8-dihydroxy-9,10-anthraquinone (XIa-2). A mixture of VIa (456 mg, 0.73 mmol) and bis(2-hydroxyethyl)amine (600 mg, 5.50 mmol) in DMF (20 mL) was stirred for 2 h and then partitioned between CHCl_3 (100 mL) and H_2O , (100 mL). The organic layer was separated, washed H_2O , 3 X 50 mL, dried (Na_2SO_4), and concentrated in vacuo. The residue was chromatographed on a silica gel column using CHCl_3 -MeOH (15:1 v/v) as the eluent. The major fraction was concentrated and the residue

dissolved in 1 N HCl. After concentration of the solution in vacuo, the residue was triturated well with MeOH (5 mL), and the crystalline HCl salt of XIa-2 (496 mg, 91%) was collected by filtration, mp 204-207 °C. Anal. ($C_{19}H_{19}NO_6HCl$) C, H, N.

5 In a similar manner but by use of the corresponding amines, XIa-1, -2, -5, -6, 8-11, and -13-15 were prepared. Also, from VIb, by the same procedure, XIb-1, -2, -5, and -6 were obtained (Table III).

EXAMPLE 17

10 3-[[N,N-Bis(2-chloroethyl)amino]methyl]-1,8-dihydroxy-9,10-anthraquinone (XIa-3). To a solution of XIa-2-HCl (1.05 g, 2.57 mmol) in dry DMF (40 mL) was added $SOCl_2$ (1.0 mL), and the solution was stirred at room temperature for 1.5 h. The solution was concentrated in vacuo (bath temperature <40°C), and the residue was cooled in an ice bath. Cold MeOH (10 mL) was added to destroy DMF-HCl complex, and the mixture was concentrated in vacuo. Upon trituration of the residue with cold MeOH (10 mL), XIa-3 (977 mg, 85.1 %) was obtained, mp 211-214 °C. Anal ($C_{19}H_{17}Cl_2NO_4HCl$) C, H, N.

20 In a similar manner but by use of the corresponding amino alcohols, IXa-3 and -7, IXb-3 and -7, Xa-3, Xb-3, XIa-7, and XIb-3 and -7 were synthesized (Table III).

25 **Spectral Studies.** Absorption spectra were measured with an IBM 9410 UV-visible spectrometer interfaced to an HP 9826 computer. Small volumes of the stock drug solutions (2 mg/mL in Me_2SO) were added to 2 mL of buffer (0.01 or 0.1 M NaCl, 5 mM Hepes, pH 7) to obtain a final drug concentration of 5-15 μM , or to the solution of native or thermally denatured DNA (0.2 and 0.1 mM, respectively) in the buffer. After incubation at room temperature for 10 min, the spectra were recorded in the 300-600 nm range (increment 1 nm) and

corrected by subtracting the spectrum of the blank which was measured before addition of the drug.

- Biological Assays. Method A.** For cell growth inhibition studies, HL-60 cells ($92.0 \times 10^5/\text{mL}$) were grown in RPMI 1640 media at 37°C in humidified 5% CO_2 for 72 h. Viable cells were counted with the trypan blue exclusion method. The fractional inhibitions at four or five concentrations of compounds (in 0.2% DMSO) were analyzed with a median-effect plot (Chou, T.-C.; et al., Adv. Enzyme Regul. 22:27 (1984)) by using a computer program. (Chou, J.; Chou, T.-C. Dose-Effect Analyses with Microcomputer. Quantitation of ID_{50} , LD_{50} , Synergism, Anagonism, Low-dose Risk, Receptor-Binding and Enzyme Kinetics; IBM-PC Series, Elsevier-Biosoft, Elsevier Scientific: Cambridge, U.K., (1986)) The median-effect concentration (ID_{50}) was automatically determined for the x intercept of the median-effect plot. Cell growth in the absence of a compound and in the presence of DMSO was used as a control. DMSO (0.2%) alone inhibited cell growth $3.8 \pm 1.2\%$ during the 72-h incubation period.
- Method B.** For precursor incorporation studies, each compound at four to six concentrations (0.2% DMSO) was preincubated with HL-60 cells ($2.5 \times 10^6/\text{Ml}$) for 15 min prior to the addition of [^3H -methyl]TdR ($1 \mu\text{Ci}$, $0.15 \text{ nmol}/\text{mL}$) and was incubated for 30 min. The incubation conditions and the procedures for isolating the DNA fractions were described previously (Chou, T.-C, et al., J. Cancer Res. 43:3074 (1983)). The incorporation of radioactivity in DNA in the absence of an analogue in the presence of DMSO was used as a control. The control value for incorporation into DNA was $8500 \pm 300 \text{ cpm}/10^6 \text{ cells}$.

EXPERIMENT 3:

- This experiment concerns Topoisomerase II (Topo II)-mediated DNA cleavage activity induced by chrysophanol derivatives.

3-(bis[2-chloroethyl]amino)methyl-1,8-dihydroxy-9,10-anthraquinone (SK-31662) and 3-(2-hydroxyethylamino)methyl-1,8-dihydroxy-9,10-anthraquinone (SK-31694) have antileukemic activity with IC_{50} s of 0.14 and 0.86 μ M, respectively. SK-31662 and 31694 inhibit kDNA decatenation at 48 and 38 μ M, respectively. The mapping of DNA-Topo II cleavage sites using Hind III-digested 3'-labeled DNA and nuclear extracts (NE) of HL-60 cells showed that at 10 μ M these two agents induce protein-linked DNA breaks with a cleavage site pattern similar to m-AMSA. They also stimulate the formation of Topo II cleavable complex in the presence of 3'-labeled DNA and NE. The amounts of protein-linked DNA induced by VP-16 and SK-31694 are reduced during 0.5 min exposure to 65 degrees C whereas SK-31662-induced protein-linked DNA complex cannot be reversed up to 15 min. The data suggest that Topo II appears to be a major cytotoxic target for these compounds and DNA intercalator with alkylating groups interact with Topo II system in an irreversible manner with enhanced toxicity.

20

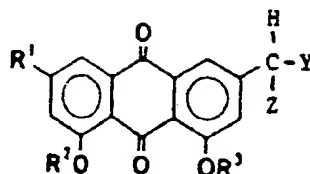
25

30

35

What is claimed is:

1. A compound having the structure:



wherein R¹ is hydrogen, a hydroxy group or a methoxy group;

R² is hydrogen or a methyl group;

R³ is hydrogen or a methyl group;

Y is a halogen, a secondary amino group or a tertiary amino group and

Z is hydrogen or a halogen.

2. A compound of claim 1, wherein the halogen is chlorine or bromine.

3. A compound of claim 1, wherein the secondary amino group comprises a lower alkyl group or a substituted lower alkyl group where each substituent is a halogen or a hydroxy, benzoxy, or acetyloxy group.

4. A compound of claim 1, wherein the tertiary amino group comprises two lower alkyl groups or substituted lower alkyl groups where each substituent is a halogen or a hydroxy, benzoxy, or acetyloxy group and where the lower alkyl groups or the substituted lower alkyl groups are the same or different.

5. A compound of claim 1 selected from the group consisting of:
- 3-Bromomethyl-1,8-dihydroxy-9,10-anthraquinone;
 - 3-Dibromomethyl-1,8-dihydroxy-9,10-anthraquinone;
 - 3-Bromomethyl-1,8-dimethoxy-9,10-anthraquinone;
 - 3-Dibromomethyl-1,8-dimethoxy-9,10-anthraquinone;
 - 3-Bromomethyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone;
 - 3-Dibromomethyl-1,8-dimethoxy-6-methoxy-9,10-anthraquinone;
 - 3-Bromomethyl-1,6,8-trimethoxy-9,10-anthraquinone; and
 - 3-Dibromomethyl-1,6,8-trimethoxy-9,10-anthraquinone.
6. A compound of claim 1 selected from the group consisting of:
- 3-(bis[2-chloroethyl]amino)methyl-1,8-dihydroxy-9,10-anthraquinone; and
 - 3-(2-hydroxyethylamino)methyl-1,8-dihydroxy-9,10-anthraquinone.
7. A compound of claim 1 selected from the group consisting of:
- 3-(N,N-Diethylamino)methyl-1,8-dihydroxy-9,10-anthraquinone;
 - 3-(N-Ethylamino)methyl-1,8-dihydroxy-9,10-anthraquinone;
 - 3-[N,N-Bis(2-hydroxyethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone;
 - 3-[N-(2-Hydroxyethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone;
 - 3-(N,N-Diethylamino)methyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone;
 - 3-(N-Ethylamino)methyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone;
 - 3-[N,N-Bis(2-hydroxyethyl)amino]methyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone;

3-[N-(2-hydroxyethyl)amino]methyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone;

3-(N,N-Diethylamino)methyl-1,8-dimethoxy-9,10-anthraquinone;

3-(N,Ethylamino)methyl-1,8-dimethoxy-9,10-anthraquinone;

3-[N,N-Bis(2-hydroxyethyl)amino]methyl-1,8-dimethoxy-9,10-anthraquinone;

3-[N-(2-Hydroxyethyl)amino]methyl-1,8-dimethoxy-9,10-anthraquinone;

3-(N,N-Diethylamino)methyl-1,6,8-trimethoxy-9,10-anthraquinone;

3-(N-Ethylamino)methyl-1,6,8-trimethoxy-9,10-anthraquinone; and

3-[N,N-Bis(2-hydroxyethyl)amino]methyl-1,6,8-trimethoxy-9,10-anthraquinone.

8. A compound of claim 1 selected from the group consisting of:

3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone;

3-[N-(2-Chloroethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone;

3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone;

3-[N-(2-Chloroethyl)amino]methyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone;

3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dimethoxy-9,10-anthraquinone;

3-[N-(2-Chloroethyl)amino]methyl-1,8-dimethoxy-9,10-anthraquinone;

3-[N,N-Bis(2-chloroethyl)amino]methyl-1,6,8-trimethoxy-9,10-anthraquinone; and

3-[N-(2-Chloroethyl)amino]methyl-1,6,8-trimethoxy-9,10-anthraquinone.

9. A pharmaceutical composition for the treatment of a malignancy in a subject comprising the compound of claim 1 or a pharmaceutically acceptable acid salt thereof and a pharmaceutically acceptable carrier, the amount of the compound or salt being 1 to 200 mg/kg of body weight of the subject.

5

10. A method of treating a subject having a malignancy which comprises administering to the subject an effective amount of the compound of claim 1 to suppress growth of the malignancy.

10

11. A method of claim 9, wherein the malignancy is leukemia.

12. A method of claim 9, wherein the malignancy is a tumor.

15

20

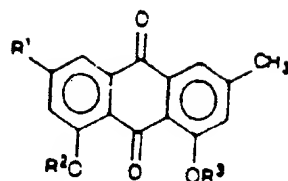
25

30

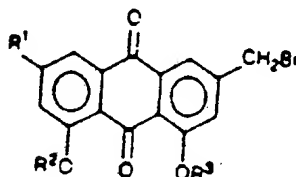
35

1/6

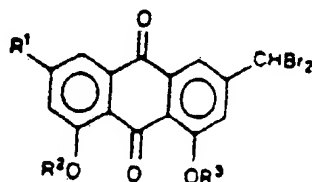
Figure 1



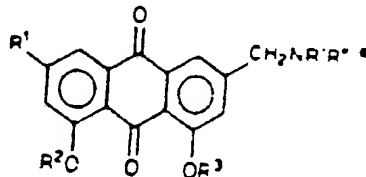
Ia $R^1, R^2, R^3 = H$ (chrysophanol)
 Ib $R^1 = OH, R^2, R^3 = H$ (modin)
 IIa $R^1 = H, R^2, R^3 = Me$
 IIb $R^1 = OMe, R^2, R^3 = Me$



IIIa $R^1 = H, R^2, R^3 = Me$
 IIIb $R^1 = OMe, R^2, R^3 = Me$
 Va $R^1 = H, R^2 = H(Me), R^3 = Me(H)$
 Vb $R^1 = OMe, R^2 = H(Me), R^3 = Me(H)$
 VIa $R^1, R^2, R^3 = H$
 VIb $R^1 = OMe, R^2, R^3 = H$



IVa $R^1 = H, R^2, R^3 = Me$
 IVb $R^1 = OMe, R^2, R^3 = Me$
 VIIa $R^1 = H, R^2 = H(Me), R^3 = Me(H)$
 VIIb $R^1 = OMe, R^2 = H(Me), R^3 = Me(H)$
 VIIIa $R^1, R^2, R^3 = H$
 VIIIb $R^1 = OMe, R^2, R^3 = H$



IXa $R^1 = H, R^2, R^3 = Me$
 IXb $R^1 = OMe, R^2, R^3 = Me$
 Xa $R^1 = H, R^2 = H(Me), R^3 = Me(H)$
 Xb $R^1 = OMe, R^2 = H(Me), R^3 = Me(H)$
 XIa $R^1, R^2, R^3 = H$
 XIb $R^1 = OMe, R^2, R^3 = H$

*See Table III for NR'R''

2/6

Figure 2

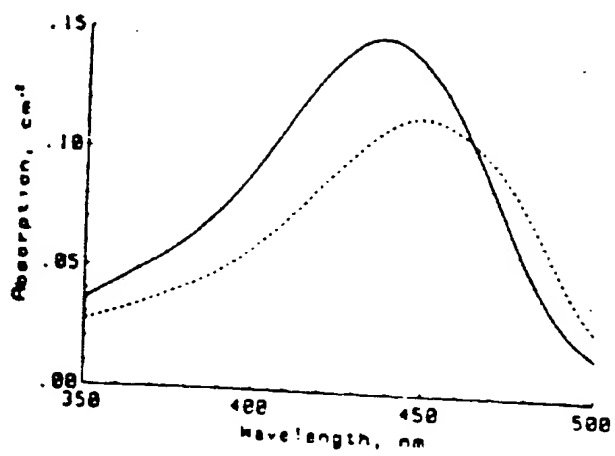
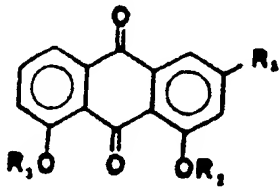
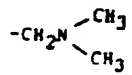
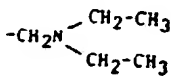
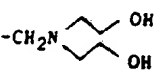
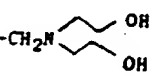
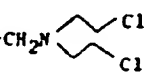


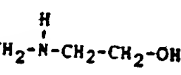
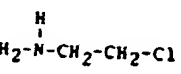
Potential Anticancer Agents

Figure 3

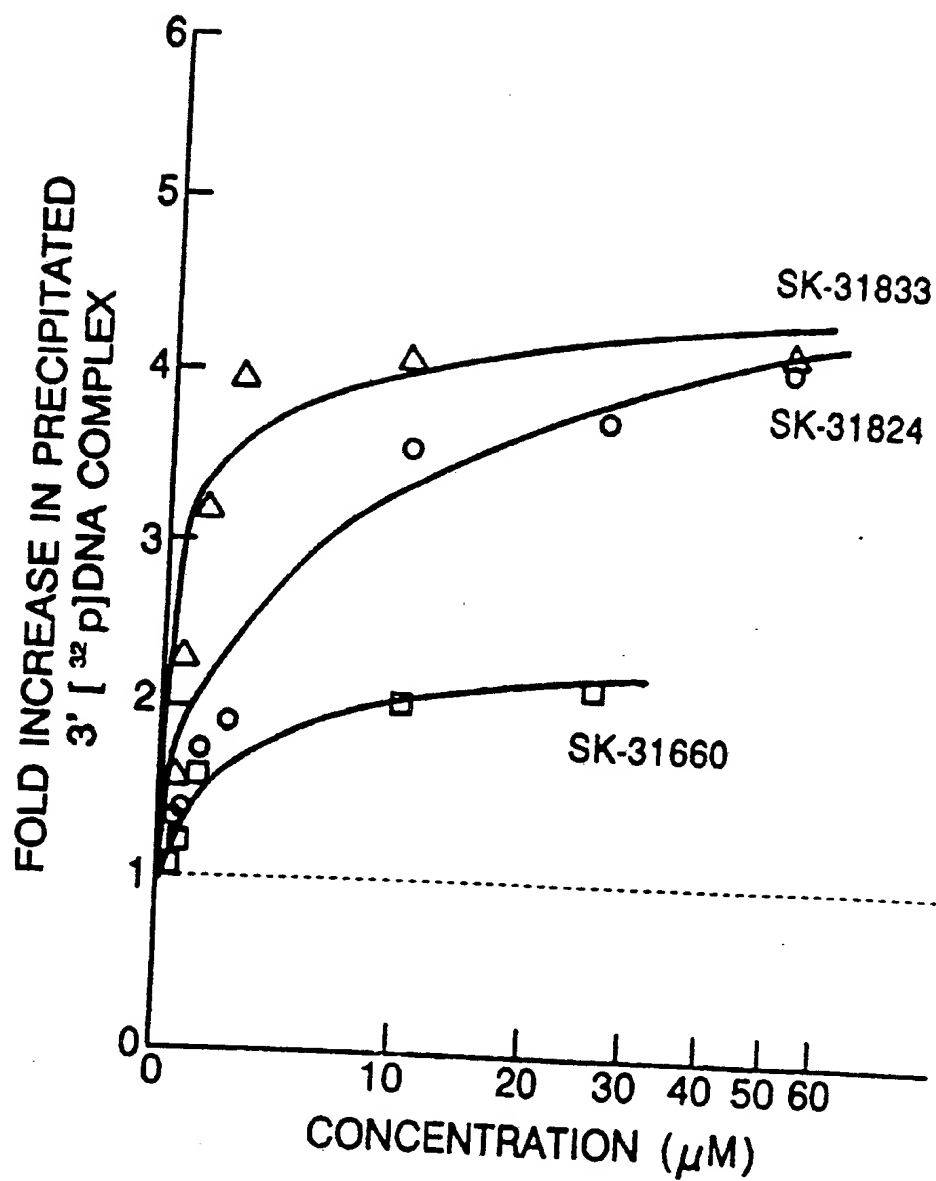
Biological Effects of Chrysophanol Derivatives

Compound				IC ₅₀ for cell growth (μ M)
	R1	R2	R3	
SK 31671		H	H	2.8
SK 31661		H	H	1.8
SK 31660		H	H	3.3
SK 31653		H	CH ₃	20.7
SK 31662		H	H	0.14
SK 31669		H	H	1.14
SK 31665		H	H	2.8
SK 31694		H	H	0.86
SK 31690		H	H	7.5
SK 31666	-CH ₂ Br	H	H	7.1

4/6

Figure 4

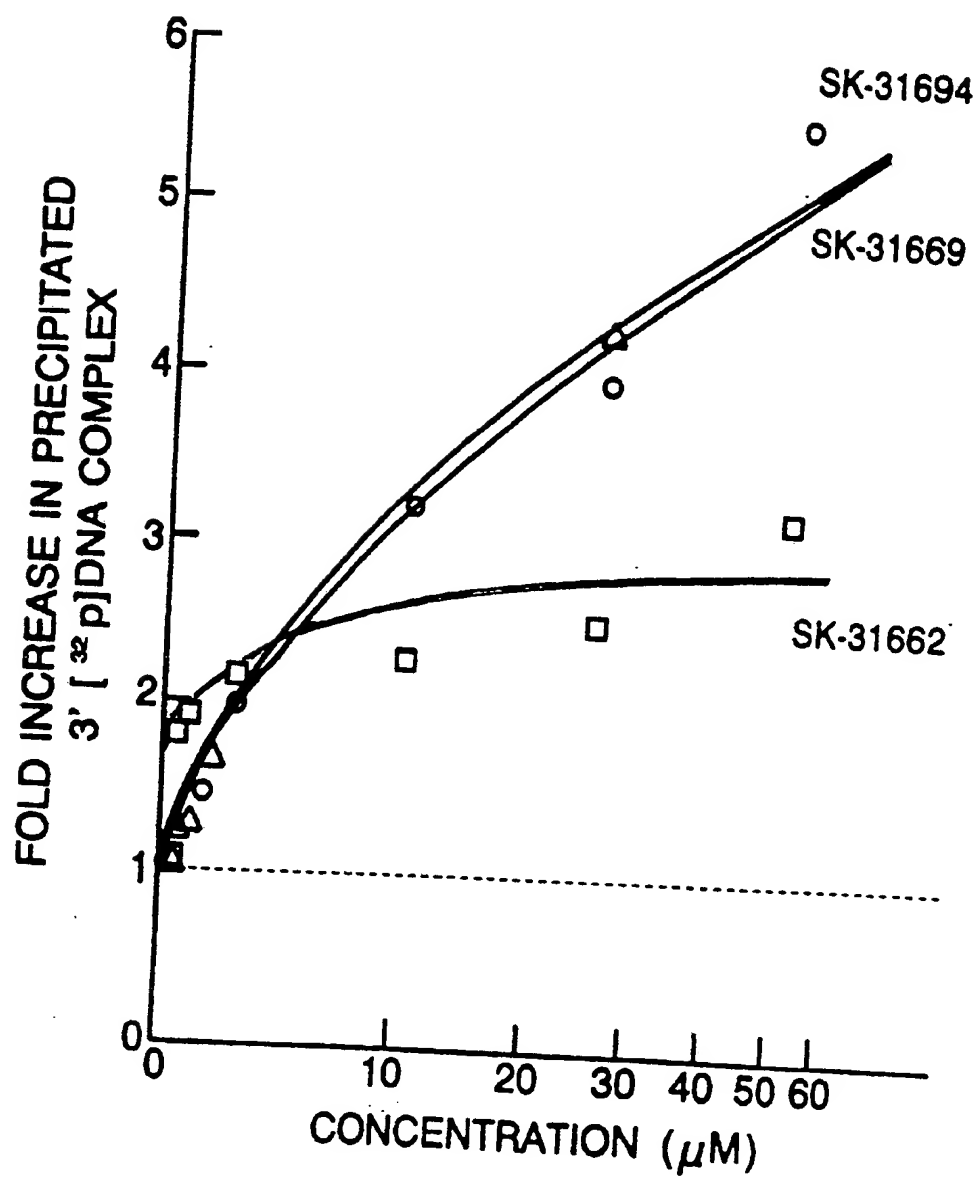
TOPO II CLEAVABLE COMPLEX FORMATION



5/6

Figure 5

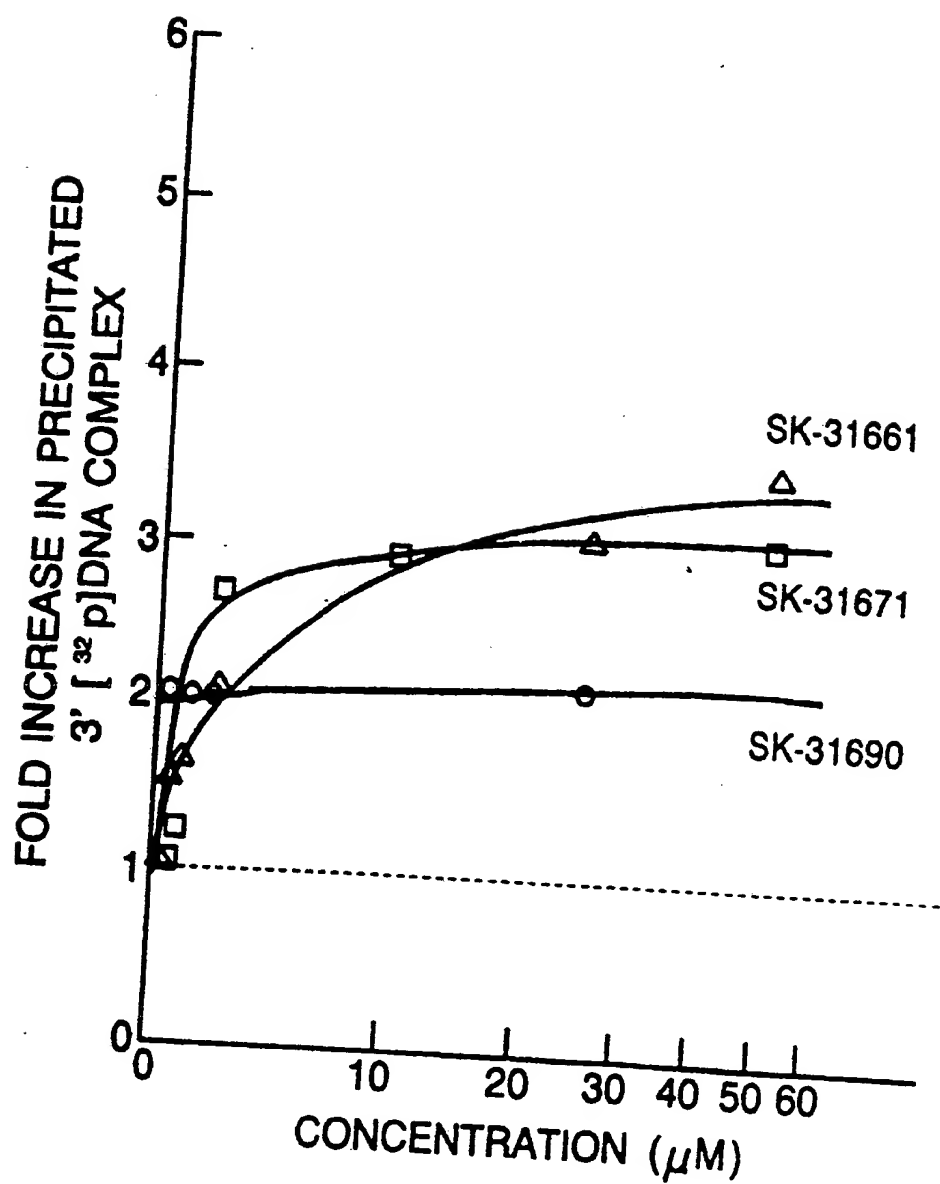
TOPO II CLEAVABLE COMPLEX FORMATION



6/6

Figure 6

TOPO II CLEAVABLE COMPLEX FORMATION



INTERNATIONAL SEARCH REPORT

International Application No. PCT/US90/00487

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) *

According to International Patent Classification (IPC) or to both National Classification and IPC
 IPC(5): C07C 50/18, 34; A61K 31/03, 09, 12, 135
 U.S. CL.: 552/261, 262, 266; 514/676, 680

II. FIELDS SEARCHED

Classification System	Minimum Documentation Searched ?
US, CL.	552/261, 262, 266; 514/676, 680

Documentation Searched other than Minimum Documentation
 to the Extent that such Documents are Included in the Fields Searched *

CAS Online: Graphic Structure

III. DOCUMENTS CONSIDERED TO BE RELEVANT :

Category *	Citation of Document, ** with indication, where appropriate, of the relevant passages **	Relevant to Claim No. ?
Y	US, A, 4,215,062 MITSCHER, 29 JULY 1980 See col. 4, lines 32-44.	1-12
Y	The Chemistry of Synthetic Dyes; Venkatoroman, "Natural Anthraquinones coloring Matters color Matters," Academic Press inc., New York, 1952 page 831.	1-12
Y	Anthracene and Anthraquinone, Barnett, "The Halogen Compounds", D VAN NOSTRAND Company, New York, 1921, pages 174-175	1-12

* Special categories of cited documents: **

"A" document defining the general state of the art which is not
 considered to be of particular relevance
 "E" earlier document but published on or after the international
 filing date
 "L" document which may throw doubts on priority claim(s) or
 which is cited to establish the publication date of another
 citation or other special reason (as specified)
 "O" document referring to an oral disclosure, use, exhibition or
 other means
 "P" document published prior to the international filing date but
 later than the priority date claimed

"T" later document published after the international filing date
 or priority date and not in conflict with the application but
 cited to understand the principle or theory underlying the
 invention
 "X" document of particular relevance; the claimed invention
 cannot be considered novel or cannot be considered to
 involve an inventive step
 "Y" document of particular relevance; the claimed invention
 cannot be considered to involve an inventive step when the
 document is combined with one or more other such docu-
 ments, such combination being obvious to a person skilled
 in the art.
 "A" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search

12 APRIL 1990

International Searching Authority

ISA/US

Date of Mailing of this International Search Report

04 MAY 1990

Signature of Authorized Officer

RAYMOND COVINGTON

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

V. ☐ OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE:

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. ☐ Claim numbers _____, because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claim numbers _____, because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out¹³, specifically:

3. ☐ Claim numbers _____, because they are dependent claims not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI. ☒ OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING:

This International Searching Authority found multiple inventions in this international application as follows:

- I. Claims 1-9 drawn to Chrysophanol products.
- II. Claims 10-12 drawn to a method for treating malignancy.

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.
2. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:
3. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:

1 - 9

4. ☐ As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority does not invite payment of any additional fee.

Remark on Protest

- ☐ The additional search fees were accompanied by applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.